Application No.: 10/586,678 Docket No.: 0599-0217PUS1
Page 3 of 9

AMENDMENTS TO THE CLAIMS

1. (Previously Presented) A fractionation device for separating one or more solutes from a raw liquid with a membrane comprising:

- 1) a supply part for loading the raw liquid;
- 2) a filtration part connected to the supply part by a flow channel for filtering out one or more solutes from the raw liquid received from the supply part to produce a filtrate;
- 3) a concentration part connected to the filtration part by a flow channel for increasing the concentration of one or more solutes in the filtrate received from the filtration part to produce a concentrated solution;
- 4) a recovery part connected to the concentration part by a flow channel for recovering the concentrated solution obtained in the concentration part; and
- 5) one or more flow pumps operatively connected to one or more of the supply part, filtration part and concentration part for moving liquid through the fractionation device, wherein the filtration part, the concentration part, and the flow channel connecting the filtration part and the concentration part form a closed circuit; and

wherein at least a portion of the circuit composed of the supply part, the filtration part, the concentration part, the recovery part, and flow channels connecting the respective parts is assembled in a cartridge,

the flow pump is a tube pump provided with a rotating rotor and a roller installed in a rotating manner in the outer circumference of the rotor, and

a portion of the outer wall of the cartridge is a squeezing member for squeezing a part of the flow channels of the circuit.

- 2. (Previously Presented) The fractionation device as claimed in claim 1, wherein the supply part, the filtration part, and the flow channel connecting the supply part and the filtration part form a closed circuit, and wherein the concentration part, the recovery part, and a flow channel connecting the concentration part and the recovery part form a closed circuit.
- 3. (Original) The fractionation device as claimed in claim 2, wherein the total inner capacity of the closed circuits is 50 mL or lower.

Application No.: 10/586,678 Docket No.: 0599-0217PUS1
Page 4 of 9

4. (Previously presented) The fractionation device as claimed in claim 2, wherein a filtration apparatus is employed in each of the filtration part and the concentration part each.

- 5. (Original) The fractionation device as claimed in claim 4, wherein the filtration apparatus is a module having hollow fiber membranes.
- 6. (Original) The fractionation device as claimed in claim 5, wherein the flow channel connecting the supply part and the filtration part is provided with a pump.
- 7. (Original) The fractionation device as claimed in claim 6, wherein the recovery part is a container for sampling a concentrated liquid.
- 8. (Original) The fractionation device as claimed in claim 7, wherein a buffer part for buffering the volumetric alteration at the time of loading the raw liquid is installed at any position in the circuit.
- 9. (Canceled).
- 10. (Canceled).
- 11. (Currently Amended) The fractionation device as claimed in claim 10 claim 1, wherein the fractionation device is provided with a transportation mechanism for transporting the cartridge in the direction to and from the rotor of the roller type tube pump to squeeze a flow pipe.
- 12. (Previously Presented) The fractionation device as claimed in claim 1, wherein the raw liquid is a body fluid or a biological component-containing solution.
- 13. (Previously Presented) A fractionation device comprising a cartridge and a roller type tube pump for separating solutes or some of the solutes in a raw liquid with a membrane,

Application No.: 10/586,678 Docket No.: 0599-0217PUS1
Page 5 of 9

wherein the cartridge comprises at least a portion of a circuit having at least a supply part for loading the raw liquid, a fractionation part connected with the supply part by a flow channel for fractionating solutes of the raw liquid by a membrane, and a recovery part connected with the fractionation part for recovering the fractionated solutes,

the circuit is a closed circuit,

a part of the outer wall of the cartridge is a squeezing member for squeezing a tube of the roller type tube pump, and

a tube forming a part of the circuit is disposed on a part of the outer wall of the squeezing member.

14. (Previously Presented) A circuit of a fractionation device for separating solutes or some of the solutes from a raw liquid with a membrane, wherein at least a portion of the circuit is contained within a cartridge and the circuit comprises a supply part for loading the raw liquid, a fractionation part connected with the supply part by a flow channel for fractionating solutes of the raw liquid with a membrane, and a recovery part connected with the fractionation part for recovering the fractionated solutes,

the circuit is a closed circuit,

a part of the outer wall of the cartridge forms a squeezing member, and

a tube forming a part of the circuit is disposed on a portion of the outer wall of the squeezing member.

- 15. (Previously Presented) A fractionation device as claimed in claim 5, wherein the module of the filtration part has a permeation ratio of human α_l microglobulin and human albumin (permeability of human α_l microglobulin/permeability of human albumin) in a range from 1.5 or higher to 1000 or lower under a condition that no antibody adsorbing proteins exist in the device, and an antibody capable of adsorbing specified proteins is contained in a middle or a rear part of the module of the filtration part.
- 16. (Previously Presented) The fractionation device as claimed in claim 15, wherein the specified proteins are serum albumin, immunoglobulin G, immunoglobulin A, immunoglobulin M, transferrin, haptoglobin, α_l -antitrypsin, α_2 -macroglobulin, α_l -acid glycoprotein, fibrinogen,

Application No.: 10/586,678 Docket No.: 0599-0217PUS1
Page 6 of 9

complement C1q, complement C3, complement C4, complement C8, complement C9, complement factor B, apolipoprotein A, apolipoprotein B, Lp(a), collagen, myosin, actin, cytokeratin, keratin, and/or fibronectin.

- 17. (Previously Presented) The fractionation device as claimed in claim 16, wherein the antibody is polyclonal antibody, monoclonal, or their fragments containing the antigen recognition sites.
- 18. (Previously Presented) The fractionation device as claimed in claim 17, wherein the antibody is fixed in the membrane surface of the module of the filtration part.
- 19. (Previously Presented) The fractionation device as claimed in claim 18, wherein the filtration part comprises columns containing hollow fiber therein and arranged in multi-step in series and the antibody is fixed in the surface in the raw liquid side of the membrane of the column in the first stage.
- 20. (Previously Presented) The fractionation device as claimed in claim 18, wherein the filtration part comprises columns containing hollow fiber therein and arranged in multi-step in series and the antibody is fixed in the surface in the permeation side of the separation membrane of the column in the first stage.
- 21. (Previously Presented) The fractionation device as claimed in claim 18, wherein the filtration part comprises columns containing hollow fiber therein and arranged in multi-step in series and the antibody exists in the mobile phase in the flow channel between the membrane of the column in a prior stage and the membrane of the column in a posterior stage.
- 22. (Previously Presented) The fractionation device as claimed in claim 18, wherein the filtration part comprises columns containing hollow fiber therein and arranged in multi-step in series and the antibody is fixed in the flow channel between the membrane of the column in a prior stage and the membrane of the column in a posterior stage.

Application No.: 10/586,678

Docket No.: 0599-0217PUS1

Page 7 of 9

23 - 27. (Cancelled)